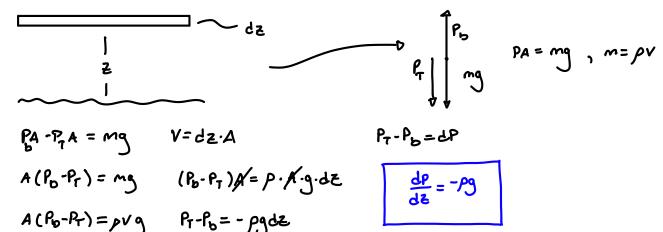
## Problem 1

a.)



b.) PY = NKT, 
$$P = \frac{Pm}{p} = NKT$$
 :  $P = \frac{Pm}{NKT}$ 

$$\frac{dP}{dz} = -g \cdot \frac{\rho_m}{N \kappa T} \qquad \therefore \qquad \int \frac{1}{\rho} d\rho = \int \frac{g_m}{N \kappa T} dz$$

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$$P(z) = ce^{-gm_{NKT}^2}$$
,  $P(z) = ce^{-gm_{NKT}^2}$ 

(c.) 
$$N_2: M_{N_2} = 28 \, \underline{9} \, , \quad O_2: M_{O_2} = 32 \, \underline{9} \, , \quad Ar: M_{Ar} = 40 \, \underline{9} \,$$

$$m_{\nu_2} = 4.65 \times 10^{-26} \frac{\text{kg}}{\text{mole}}$$
,  $m_{e_2} = 5.32 \times 10^{-26} \frac{\text{kg}}{\text{mole}}$ ,  $m_{Ar} = 6.64 \times 10^{-26} \frac{\text{kg}}{\text{mole}}$ 

(e.) 
$$p(z=0) = \frac{1.01 \times 10^5 \text{ Pa} \cdot 4.81 \times 10^{-36} \text{ Mg/molecule}}{1.38 \times 10^{-33} \text{ J/k} \cdot 293 \text{ K}} = 1.2 \text{ Mg}$$

$$\frac{p \cdot n \kappa \tau}{m} = ce^{-gm2/n \kappa \tau} : \frac{p_N \kappa \tau}{cm} = e^{-gm2/n \kappa \tau} \cdot \ln \left(\frac{p_N \kappa \tau}{cm}\right) = \frac{-gm2}{n \kappa \tau}$$

$$Z = -ln\left(\frac{p_{NKT}}{cm}\right) \cdot \frac{NKT}{gm} = 19,761.7 m$$

Z= 19,762 m

## Problem 2

Constant Volume, pressure doubled

$$\left( \begin{array}{c} P + \frac{N^2}{V^2} \alpha \end{array} \right) \left( \begin{array}{c} V - Nb \end{array} \right) = NKT : \frac{N^2}{V^2} \alpha \longrightarrow \alpha \quad V - nb \longrightarrow \beta \quad NK \longrightarrow \delta$$

$$(P_{i} + \omega)(\beta) = x \cdot T_{i} : T_{i} = (P_{i} + \omega)(\beta) : (2P_{i} + \omega)(\beta) = x \cdot T_{f}$$

$$P_{i} = \frac{x \cdot T_{i}}{\beta} - \omega : (2 \cdot (\frac{x \cdot T_{i}}{\beta} - \omega) + \omega)(\beta) = x \cdot T_{f}$$

$$(\frac{2x \cdot T_{i}}{\beta} - \omega)(\beta) = x \cdot T_{f}$$

$$(\frac{2x \cdot T_{i}}{\beta} - \omega)(\beta) = x \cdot T_{f}$$

$$(2x \cdot T_{i} - \omega)(\beta) = x \cdot T_{f} : T_{f} = 2T_{i} - \omega\beta$$

Tf = 
$$\partial T_i - \alpha B$$
, .. The temperature increases by less than a factor of two. iii.)

## Problem 3

$$\left(P + \frac{N^2}{v^2} a\right) \left(V - Nb\right) = NKT$$
,  $N_A = N_B$ ,  $T_A = T_B$ ,  $V_A = V_B$ 

a.) The only thing that could cause gos A to have a greater pressure than gas B of Vice versa are the coefficients: a and b in the van der Waals gas equation. More precisely, if attractive forces of the molecules in gas A are greater than that of gas B, then we should expect the pressure of B to be greater than that of gas A.

Therefore we shouldn't expect the pressures to be the same

- b.) The ideal gas law predicts the pressure of these gases to be the same. This could be correct if the coefficient of a and b were the same for each gas.
- C.) The van der wasses equation of state could predict a difference in pressure of the gases if and only if the "a" coefficient were different for each gas. For instance if b were the same for each gas, then

if aas ab then Pb>Pa
if aas ab then Pb<Pa

and thus van der Woods equation State will predict different pressures of these gases.